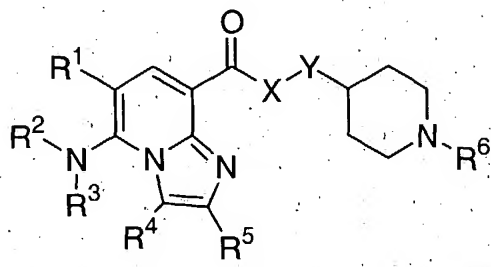


AMENDMENTS TO THE CLAIMS

1. - 7. (canceled)

8. (currently amended) A pharmaceutical composition for the ~~treatment or~~ prevention of disease conditions mediated by 5-HT₄ receptor activity, in a mammalian subject, which comprises a therapeutically effective amount of a ~~compound of Claim 1~~ a compound of the formula (I):



(I)

or the pharmaceutically acceptable salts thereof wherein

R¹ is hydrogen, halo or C₁₋₆ alkyl;

R² and R³ are independently hydrogen, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, mono- or di-(C₁₋₅)alkyl amino, amino(C₁₋₅)alkyl or hydroxy(C₁₋₅)alkyl; or R² and R³ taken together with the nitrogen atom to which they are attached may form substituted or non-substituted nitrogen-containing heterocyclic;

R⁴ is hydrogen, halo, C₁₋₈ acyl, amino, amido, substituted or non-substituted aryl, substituted or non-substituted aryl(C₁₋₆)alkyl; or substituted or non-substituted heterocyclic;

R⁵ is hydrogen, halo, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₁₋₈ acyl, amino, amido, substituted or non-substituted aryl, substituted or non-substituted aryl(C₁₋₆)alkyl, or substituted or non-substituted heterocyclic;

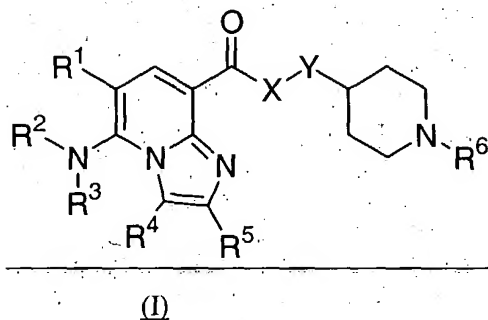
R⁶ is hydrogen, C₁₋₆ alkyl or C₁₋₆ alkoxy (C₁₋₆)alkyl;

X is NR⁹ wherein R⁹ is hydrogen or C₁₋₆ alkyl; and

Y is (CR⁷R⁸)_n wherein R⁷ and R⁸ are independently hydrogen or C₁₋₆ alkyl, and n is an integer from 0 to 5;

and a pharmaceutically acceptable carrier.

9. (currently amended) A pharmaceutical composition for the ~~treatment or~~ prevention of gastroesophageal reflux disease, gastrointestinal disease, gastric motility disorder, upper gut motility disorder, non-ulcer dyspepsia, Functional dyspepsia, irritable bowel syndrome, constipation, dyspepsia, esophagitis, gastroesophageal disease, ~~nausea~~ nausea, central nervous system disease, alzheimers disease, cognitive disorder, emesis, migraine, neurological disease, pain, ischaemic stroke, anxiety or cardiovascular disorder, which comprises a therapeutically effective amount of ~~a compound of Claim 1~~ a compound of the formula (I):



or the pharmaceutically acceptable salts thereof wherein

R¹ is hydrogen, halo or C₁₋₆ alkyl;

R² and R³ are independently hydrogen, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, mono- or di-(C₁₋₅)alkyl amino, amino(C₁₋₅)alkyl or hydroxy(C₁₋₅)alkyl; or R² and R³ taken together with the nitrogen atom to which they are attached may form substituted or non-substituted nitrogen-containing heterocyclic;

R⁴ is hydrogen, halo, C₁₋₈ acyl, amino, amido, substituted or non-substituted aryl, substituted or non-substituted aryl(C₁₋₆)alkyl, or substituted or non-substituted heterocyclic;

R⁵ is hydrogen, halo, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₁₋₈ acyl, amino, amido, substituted or non-substituted aryl, substituted or non-substituted aryl(C₁₋₆)alkyl, or substituted or non-substituted heterocyclic;

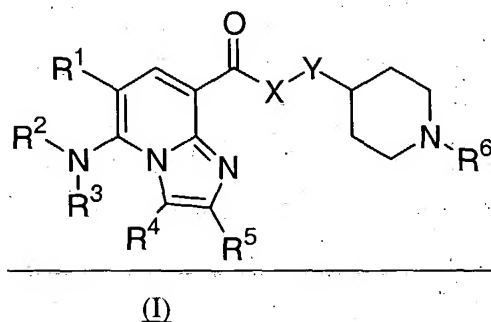
R⁶ is hydrogen, C₁₋₆ alkyl or C₁₋₆ alkoxy (C₁₋₆)alkyl;

X is NR⁹ wherein R⁹ is hydrogen or C₁₋₆ alkyl; and

Y is $(CR^7R^8)_n$ wherein R^7 and R^8 are independently hydrogen or C_{1-6} alkyl, and n is an integer from 0 to 5;

and a pharmaceutically acceptable carrier.

10. (currently amended) A method for the ~~treatment or~~ prevention of disease conditions mediated by 5-HT₄ receptor activity, in a mammalian subject, which comprises administering to said subject a therapeutically effective amount of a ~~compound according to Claim 1~~ a compound of the formula (I):



or the pharmaceutically acceptable salts thereof wherein

R^1 is hydrogen, halo or C_{1-6} alkyl;

R^2 and R^3 are independently hydrogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, mono- or di-
(C_{1-5})alkyl amino, amino(C_{1-5})alkyl or hydroxy(C_{1-5})alkyl; or R^2 and R^3 taken together with
the nitrogen atom to which they are attached may form substituted or non-substituted
nitrogen-containing heterocyclic;

R^4 is hydrogen, halo, C_{1-8} acyl, amino, amido, substituted or non-substituted aryl, substituted
or non-substituted aryl(C_{1-6})alkyl, or substituted or non-substituted heterocyclic;

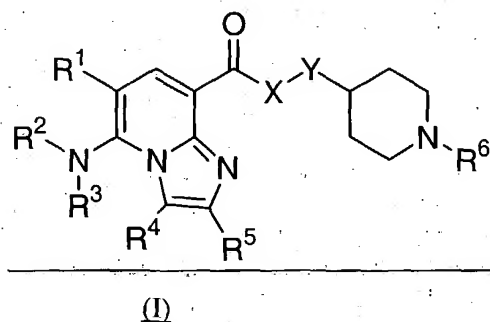
R^5 is hydrogen, halo, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-8} acyl, amino, amido,
substituted or non-substituted aryl, substituted or non-substituted aryl(C_{1-6})alkyl, or
substituted or non-substituted heterocyclic;

R^6 is hydrogen, C_{1-6} alkyl or C_{1-6} alkoxy (C_{1-6})alkyl;

X is NR^9 wherein R^9 is hydrogen or C_{1-6} alkyl; and

Y is $(CR^7R^8)_n$ wherein R^7 and R^8 are independently hydrogen or C_{1-6} alkyl, and n is an
integer from 0 to 5.

11. (currently amended) A method for the ~~treatment or~~ prevention of gastroesophageal reflux disease, gastrointestinal disease, gastric motility disorder, upper gut motility disorder, non-ulcer dyspepsia, Functional dyspepsia, irritable bowel syndrome, constipation, dyspepsia, esophagitis, gastroesophageal disease, ~~aa~~nausea, central nervous system disease, alzheimers disease, cognitive disorder, emesis, migraine, neurological disease, pain, ischaemic stroke, anxiety or cardiovascular disorder, which comprises administering to said subject a therapeutically effective amount of ~~a compound according to Claim 1~~ a compound of the formula (I):



or the pharmaceutically acceptable salts thereof wherein

R¹ is hydrogen, halo or C₁₋₆ alkyl;

R² and R³ are independently hydrogen, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, mono- or di-(C₁₋₅)alkyl amino, amino(C₁₋₅)alkyl or hydroxy(C₁₋₅)alkyl; or R² and R³ taken together with the nitrogen atom to which they are attached may form substituted or non-substituted nitrogen-containing hetrocyclic;

R⁴ is hydrogen, halo, C₁₋₈ acyl, amino, amido, substituted or non-substituted aryl, substituted or non-substituted aryl(C₁₋₆)alkyl, or substituted or non-substituted heterocyclic;

R⁵ is hydrogen, halo, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₁₋₈ acyl, amino, amido, substituted or non-substituted aryl, substituted or non-substituted aryl(C₁₋₆)alkyl, or substituted or non-substituted heterocyclic;

R⁶ is hydrogen, C₁₋₆ alkyl or C₁₋₆ alkoxy (C₁₋₆)alkyl;

X is NR⁹ wherein R⁹ is hydrogen or C₁₋₆ alkyl; and

Y is $(CR^7R^8)_n$ wherein R^7 and R^8 are independently hydrogen or C_{1-6} alkyl, and n is an integer from 0 to 5.

12. – 13. (canceled)